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CLAIMS:

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- 1. A C8-substituted purine nucleotide analog, wherein the analog is substituted at the C8 position with a substituent other than H.
- 2. The analog of claim 1, wherein the purine is adenine.
- 3. The analog of claim 1, wherein the substituent is an 10 ether, thioether or an amine.
 - 4. The analog of claim 2, wherein the substituent is an ether, thioether or an amine.
- 15 5. The analog of claim 1, wherein the substituent is an ether, and wherein the ether substituent has the structure:
 - 6. The analog of claim 5, wherein X is an alkyl group.

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- 7. The analog of claim 6, wherein X is selected from the group consisting of:
 - (a) C₇H₁₃ (cycloheptyl)
 - (b) $(CH_3)_3CCH_2$
- 25 (c) $CH_3(CH_2)_n$, wherein lsns5

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The analog of claim 1, wherein the substituent is a thioether, and wherein the thioether substituent has the structure.

S-X.

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9. The analog of claim 8, wherein X is an alkyl group.

10. The analog of claim 9, wherein X is selected from the group consisting of:

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- (a) \setminus C₇H₁₃ (cycloheptyl)
- (b) \ (CH₃)₃CCH₂
- (c) \backslash CH₃ (CH₂)_n, wherein 1sns5

11. The analog of claim 1, wherein the substituent is an 15 amine, and wherein the amine substituent has the structure:
-NH-X.

12. The analog of claim 11, wherein X is an alkyl group.

20 13. The analog of claim 12, wherein X is selected from the group consisting of:

- (a) C_7H_{13} (cycloheptyl)
- (b) (CH₃)₃CCH₂
- (c) $CH_3(CH_2)_n$, wherein $1 \le n \le 5$

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The analog of claim 1 selected from the group consisting of:

compound 6a, compound 6b, compound 6c, compound 6d, compound 6e, compound 7a, compound 7b, compound 7c, compound 7d, compound 7e, compound 8a, compound 8b, compound &c, compound &d, and compound &e.

A method for modulating the activity of an NTPDase 15. enzyme comprising exposing the enzyme to the analog according 10 to claim 1.

The method according to claim 15 wherein the activity 16. of the NTPDase enzyme is inhibited.

A method for modulating the level of purine 17. nucleotide(s) and/or nucleoside(s) and/or metabolite(s) or derivative(s) thereof in a biological system, comprising the step of introducing into said system the analog according to claim 1.

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A method for modulating the activity of a biological 18. process in a biological system, wherein said process is affected by the level ϕf purine nucleotide(s) and/or nucleoside(s) and/or medabolite(s) or derivative(s) thereof in 25 said system, comprising the step of introducing into said system the analog according to claim 1.

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19. The method of claim 18, wherein the biological process is aggregation and thrombogenecity.

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